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UTILITY PATENT APPLICATION TRANSMITTAL (Large Entity)

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Docket No.
TAM-1629DIV2

Total Pages in this Submission

TO THE ASSISTANT COMMISSIONER FOR PATENTS

Box Patent Application
Washington, D.C. 20231

Transmitted herewith for filing under 35 U.S.C. 111(a) and 37 C.F.R. 1.53(b) is a new utility patent application for an invention entitled:

SUBSTRATE TREATMENT METHOD

and invented by:

Leverenz, et al.

If a CONTINUATION APPLICATION, check appropriate box and supply the requisite information:

☐ Continuation ☒ Divisional ☐ Continuation-in-part (CIP) of prior application No.: 09/095,398

Which is a:

☐ Continuation ☐ Divisional ☐ Continuation-in-part (CIP) of prior application No.:

Which is a:

☐ Continuation ☐ Divisional ☐ Continuation-in-part (CIP) of prior application No.:

Enclosed are:

Application Elements

1. ☒ Filing fee as calculated and transmitted as described below
2. ☒ Specification having _____ pages and including the following:
 - a. ☒ Descriptive Title of the Invention
 - b. ☐ Cross References to Related Applications (if applicable)
 - c. ☐ Statement Regarding Federally-sponsored Research/Development (if applicable)
 - d. ☐ Reference to Microfiche Appendix (if applicable)
 - e. ☒ Background of the Invention
 - f. ☒ Brief Summary of the Invention
 - g. ☒ Brief Description of the Drawings (if drawings filed)
 - h. ☒ Detailed Description
 - i. ☒ Claim(s) as Classified Below
 - j. ☒ Abstract of the Disclosure

JC877 U.S. PTO
09/627801

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UTILITY PATENT APPLICATION TRANSMITTAL (Large Entity)

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Docket No.
TAM-1629DIV1

Total Pages in this Submission

Application Elements (Continued)

3. ☒ Drawing(s) (when necessary as prescribed by 35 USC 113)
- a. ☐ Formal Number of Sheets _____
- b. ☒ Informal Number of Sheets 11
4. ☒ Oath or Declaration
- a. ☐ Newly executed (original or copy) ☐ Unexecuted
- b. ☒ Copy from a prior application (37 CFR 1.63(d)) (for continuation/divisional application only)
- c. ☒ With Power of Attorney ☐ Without Power of Attorney
- d. ☐ DELETION OF INVENTOR(S)
Signed statement attached deleting inventor(s) named in the prior application,
see 37 C.F.R. 1.63(d)(2) and 1.33(b).
5. ☒ Incorporation By Reference (usable if Box 4b is checked)
The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.
6. ☐ Computer Program in Microfiche (Appendix)
7. ☐ Nucleotide and/or Amino Acid Sequence Submission (if applicable, all must be included)
- a. ☐ Paper Copy
- b. ☐ Computer Readable Copy (identical to computer copy)
- c. ☐ Statement Verifying Identical Paper and Computer Readable Copy

Accompanying Application Parts

8. ☐ Assignment Papers (cover sheet & document(s))
9. ☐ 37 CFR 3.73(B) Statement (when there is an assignee)
10. ☐ English Translation Document (if applicable)
11. ☒ Information Disclosure Statement/PTO-1449 ☒ Copies of IDS Citations
12. ☒ Preliminary Amendment
13. ☒ Acknowledgment postcard
14. ☒ Certificate of Mailing
- ☐ First Class ☒ Express Mail (Specify Label No.): _____

UTILITY PATENT APPLICATION TRANSMITTAL
(Large Entity)

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Docket No.
TAM-1629DIV2

Total Pages in this Submission

Accompanying Application Parts (Continued)

15. ☐ Certified Copy of Priority Document(s) (if foreign priority is claimed)

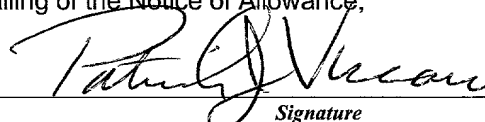
16. ☐ Additional Enclosures (please identify below):

Fee Calculation and Transmittal

CLAIMS AS FILED

For	#Filed	#Allowed	#Extra	Rate	Fee
Total Claims	11	- 20 =	0	x \$18.00	\$0.00
Indep. Claims		- 3 =	0	x \$78.00	\$0.00
Multiple Dependent Claims (check if applicable) <input type="checkbox"/>					\$0.00
BASIC FEE					\$690.00
OTHER FEE (specify purpose)					\$0.00
TOTAL FILING FEE					\$690.00

- ☐ A check in the amount of _____ to cover the filing fee is enclosed.
- ☒ The Commissioner is hereby authorized to charge and credit Deposit Account No. **01-0840** as described below. A duplicate copy of this sheet is enclosed.
- ☒ Charge the amount of **\$690.00** as filing fee.
- ☒ Credit any overpayment.
- ☒ Charge any additional filing fees required under 37 C.F.R. 1.16 and 1.17.
- ☐ Charge the issue fee set in 37 C.F.R. 1.18 at the mailing of the Notice of Allowance, pursuant to 37 C.F.R. 1.311(b).


Signature

Patrick J. Viccaro, Reg. No. 27,842
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Dated: July 28, 2000

CC:

Attorney Docket No. TAM-1629DIV2

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of

Roy V. Leverenz, et al.

TITLE: SUBSTRATE TREATMENT
METHOD

PRELIMINARY AMENDMENT

Pittsburgh, Pennsylvania 15222

July 28, 2000

Hon. Commissioner of Patents
and Trademarks
Washington, D.C. 20231

Sir:

Applicants respectfully request entry of the following
amendments in the above-identified application prior to
examination of the application:

In The Claims:

Delete claims 1 through 29 and 31 through 38. Claims 30
and 39 through 48 remain.

In the Specification:

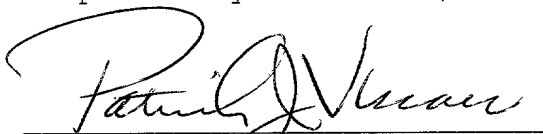
On page 1, amend the Title of this application to read
"COATED SUBSTRATE". On page 1, after Title, insert phrase "This
is a divisional application of U.S. 09/095,398, filed June 10,
1998".

09/095,398 "COATED SUBSTRATE"

REMARKS

The present invention is a Division Application of U.S. 09/095,398 filed June 10, 1998. It is respectfully submitted that the claims of the present application, as amended herein, should be held patentable, and passage of the claims to allowance at an early date is respectfully requested. Should the Examiner have any concerns after considering the present Preliminary Amendment, he or she is invited to contact the undersigned at the telephone number listed below so that those concerns may be expeditiously addressed.

Respectfully submitted,


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INVENTORS

Roy V. Leverenz and John Bost

TITLE

5 Substrate Treatment Method

TECHNICAL FIELD AND INDUSTRIAL APPLICABILITY OF THE INVENTION

10 The present invention relates to a method for etching composite material
substrates and other substrates, and also is directed to a method for applying wear-
resistant and other coatings to composite material substrates and other substrates. The
present invention also relates to composite material substrates, which are comprised of
particles of a hard constituent phase in a binder material phase that binds together the
15 hard constituent particles, having wear-resistant and other coatings. The present
invention finds application in any field in which it is advantageous to enhance the
adhesion of a wear-resistant and other types of coatings to substrates. Examples of
fields of application of the present invention include the manufacture and treatment of
dies used in metal stamping, punching, threading, and blanking, and the manufacture
20 and treatment of metal cutting inserts used in milling, turning, drilling, boring, and
other metal removal operations.

BACKGROUND OF THE INVENTION

Composite materials comprised of particles of a hard constituent phase and a
25 binder phase binding the particles together are common and are referred to as

“composite materials” or “composite substrates” hereinafter. Such materials also may be referred to as “cemented” composite materials and include, for example, ceramics, cermets, and cemented carbides. Cemented carbides, include, for example, materials composed of a hard particulate material such as, for example, particles of one or more of tungsten carbide (WC), titanium carbide (TiC), titanium carbonitride (TiCN), tantalum carbide (TaC), tantalum nitride (TaN), niobium carbide (NbC), niobium nitride (NbN), zirconium carbide (ZrC), zirconium nitride (ZrN), hafnium carbide (HfC), and hafnium nitride (HfN) cemented together by a binder phase that is composed predominantly of one or more of cobalt, nickel, and iron.

10 Metal cutting inserts fabricated from composite materials are commonly used in chip cutting machining of metals in the metal machining industry. Metal cutting inserts are commonly fabricated from particles of metal carbide, usually tungsten carbide with the addition of carbides of other metals such as, for example, niobium, titanium, tantalum, and a metallic binder phase of cobalt or nickel. The carbide materials provide high strength but still may wear quickly when used in, for example, 15 milling and other metal machining operations. By depositing a thin layer of wear-resistant material on the working surfaces of cemented carbide cutting inserts it is possible to increase the wear-resistance of the inserts without adversely affecting toughness. Commonly used wear-resistant cemented carbide insert coatings include, 20 for example, TiC, TiN, TiCN, and Al_2O_3 . Such wear-resistant coatings reduce the erosion and corrosion of the inserts' binder material.

The utility of coated composite materials such as coated cemented carbides is limited by the strength of adhesion of the wear-resistant coating to the composite

material. Absence of strong adhesion between wear-resistant coatings and metal cutting inserts causes delamination of the coatings from the inserts, decreasing the inserts' service life. The presence of cobalt at the inserts' surfaces also increases the tendency of the coatings and substrates to experience delamination during use.

- 5 Accordingly, it would be advantageous to provide a novel method for increasing the adhesion of wear-resistant coatings to composite materials. More broadly, it would be advantageous to enhance the adhesion of wear-resistant coatings and other types of coatings to composite material and other types of substrates.

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SUMMARY OF THE INVENTION

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The present invention provides a method for removing a portion of the binder phase from a substrate that is composed of at least particles of a first phase joined together by the binder phase. The present method includes the step of etching at least a portion of a surface of the substrate by contacting the surface with a gas flow that is composed of at least an etchant gas and a second gas for a time period that will allow for removal of the desired amount of binder phase. The second gas comprises one or more gases that will not react with the substrate or the removed binder material and that will not alter the oxidation state of the substrate during the etching step. Preferably, the second gas is one or more gases that will not react with the substrate or the removed portion of binder material to form deposits of a phase of W_xCo_yC (wherein $x = 3-9$ and $y = 2-6$), also referred to herein as an η (eta) phase, on the substrate.

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The etchant gas used in the present method may be any gas or combination of gases that will suitably remove the desired portion of the binder phase from the substrate during the etching step. Possible etchant gases include hydrogen chloride gas, H_2F_2 gas, and gaseous forms of any of the Group VIIA elements. Other possible etchant gases useful in the present method will be apparent to those having ordinary skill once apprised of the present invention. The second gas may be, for example, one or more gases selected from nitrogen gas, helium gas, argon gas, and neon gas. Preferably the gas flow is applied to the substrate during the etching step by introducing a flow of the etchant gas concurrently with a flow of the second gas into a chamber containing the substrate at a pressure and temperature, and for a time, that will result in removal of the desired portion of the binder phase. In one particular application of the present method, the gas flow consists of concurrent flows of hydrogen chloride gas and nitrogen gas.

Preferably, during the etching step binder phase is removed from a surface of the substrate to a depth of between about 3 microns to about 15 microns, and more preferably to a depth of between about 4 microns to about 6 microns.

The method of the present invention preferably is applied to substrates composed of a composite material comprising particles of a hard constituent material joined together by a binder material. Examples of such composite materials include cemented carbides and cermets. Examples of the binder material of such composite materials include materials composed of one or more materials selected from cobalt, nickel, iron, elements within Group VIII of the periodic table, copper, tungsten, zinc, and rhenium. Once apprised of the details of the present invention, one of ordinary

skill in the substrate coating and treatment arts will comprehend additional composite materials to which the present invention may be applied.

The present invention also is directed to a method for applying a coating to at least a portion of the surface of a substrate, preferably a composite substrate that includes hard constituent material particles joined together by a binder. The method is carried out by removing a portion of the binder from a surface of the substrate by contacting the surface with a gas flow including an etchant gas and a second gas for a period of time that will remove the desired portion of binder. The surface etching effect of the etchant gas provides an etched surface on the substrate, and the etched surface will include voids produced as the binder is etched away from between hard constituent particles. The second gas is one or more gases that will not react with the substrate or the portion of the binder removed from the substrate, and that will not change the oxidation state of the substrate during the etching process. Preferably, the second gas will not react during the etching process to form eta phase within the voids etched in the substrate's surface. In a subsequent step of the method, a coating is applied to at least a portion of the etched surface. At least a portion of the coating is deposited within at least a portion of voids on the etched surface created by removal.

Thus, the etching step of the present invention may be preceded or followed by one or more additional steps, including, for example, the step of depositing a coating on the etched surface of the substrate produced by the etching step. Because the coating infiltrates voids in the etched surface of the substrate that have been produced by removal of binder material during the etching step, the adhesion of the coating to the substrate is enhanced. Preferably, the coating is one that enhances the wear

resistance of the substrate, but it also may be selected from any other conventional substrate coating. Possible wear-resistant coatings that may be applied in the coating step of the present method include those composed of, for example, one or more of TiC, TiN, TiCN, diamond, Al₂O₃, MT-milling coating (described in detail below),
5 TiAlN, HfN, HfCN, HfC, ZrN, ZrC, ZrCN, BC, Ti₂B, MoS, Cr₃C₂, CrN, CrCN, and CN.

The present invention is also directed to substrates that have been produced by the method of the present invention. For example, such substrates within the scope of the invention may have an etched surface produced by the foregoing etching step, and
10 also may have a coating, wear-resistant or otherwise, which at least partially infiltrates voids produced in the substrate's surface by the etching step of the invention. In particular, the present invention is directed to a substrate composed of a composite material including particles of a hard constituent material and a binder material. The substrate includes an etched surface portion having voids thereon produced by
15 removing a portion of the binder material therefrom by contacting the surface portion with concurrent flows of at least a suitable etchant gas and a second gas. The second gas must be incapable of reacting with the substrate or the removed binder material or changing the oxidation state of the substrate during etching of the binder material. A coating may be adhered to at least a portion of the etched surface portion of the
20 substrate, and at least a portion of the coating is deposited within at least a portion of the voids provided in the etched surface portion.

Examples of applications of the method of the present invention include the manufacture and treatment of wear resistant cutting inserts, dies, punches, and other

elements used in applications such as: metal stamping, punching, threading, blanking, milling, turning, drilling, boring, and other metal removal operations; mining and oil drilling, including fabricating or treating mining and drilling bits used in long wall and coal boring miners, tricone, percussive and rooftop drilling bits, road planing and other like applications; wood working applications, including fabricating or treating bits and blades used in sawing, planing, routing, shaping, and other woodworking applications; drawing, heading, and back extrusion, including the fabrication and treatment of punches and dies used in such applications; rod mill rolls; and high corrosion environments. An example of a specific application of the present invention is in the manufacture and treatment of items made from tungsten-based alloys containing iron, nickel, copper and/or cobalt. Such items include, for example, aircraft weights, electrical contact points, and electrodes.

The reader will appreciate the foregoing details and advantages of the present invention, as well as others, upon consideration of the following detailed description of the invention. The reader also may comprehend such additional details and advantages of the present invention upon practicing the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

The features and advantages of the present invention may be better understood by reference to the accompanying figures, in which:

Figure 1 is a photomicrograph of a prepared section of a metal cutting insert composed of SD-5 material coated with wear-resistant MT-milling (moderate temperature milling and turning) coating by the method of the present invention;

Figures 2a-2c and 3a-3c are photomicrographs showing the condition of an edge surface of each of three metal cutting inserts, composed of SD-5 material and coated with an MT-milling coating by the method of the invention, after 10 and 18 milling passes, respectively;

5 Figures 4a-4c and 5a-5c are photomicrographs showing the condition of an edge surface of each of three uncoated metal cutting inserts, composed of SD-5 material, after 10 and 18 milling passes, respectively;

Figures 6a-6c are photomicrographs showing the condition of an edge surface of three metal cutting inserts, composed of T-14 material and coated with an
10 MT-milling coating by the method of the invention, after 4 milling passes;

Figures 7a-7c are photomicrographs showing the condition of an edge surface of three metal cutting inserts composed of T-14 material, each insert both unetched and uncoated, after 4 milling passes;

Figure 8 is a photomicrograph of a metal cutting insert composed of H-91
15 material and coated with an MT-milling coating by the method of the present invention;

Figures 9 is a photomicrograph showing the condition of a metal cutting insert, composed of H-91 material and coated with an MT-milling coating by the method of the invention, after one milling pass;

20 Figure 10 is a photomicrograph showing the condition of a metal cutting insert, composed of H-91 material and coated with an MT-milling coating of TiN/TiCN/TiN layers totaling approximately 5 microns (applied by CVD), after one milling pass; and

Figures 11a-11d are photomicrographs of a prepared section of a heavy metal part containing tungsten metal particles (about 90 weight percent of the part's total weight) suspended in an iron/nickel binder (about 10 weight percent of the part's total weight) that was etched and coated with an MT-milling coating by the method of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

An aspect of the present invention is directed to a method for applying a coating, preferably a wear-resistant coating, to a composite material substrate. The composite material substrate includes a phase of a hard constituent and also includes a binder phase that is predominantly one or more of cobalt, nickel, and iron. The present inventors have discovered that the method of the invention enhances the adherence of the coating to the composite material substrate and inhibits delamination of the coating. The present invention also is directed to etched and etched/coated substrates prepared by the method of the present invention.

It is believed that in relation to the known composite material substrate coating methods, the present method improves adhesion between composite material substrates and wear-resistant coatings by allowing the coatings to infiltrate the surface of the substrate. To accomplish this, a portion of the binder phase of a surface region of the composite material substrate is removed by a novel etching procedure, preferably to a depth in the range of about 3 to about 15 microns (inclusive), while leaving the hard constituent particles in the surface region substantially intact. Wear-resistant coatings applied to composite material substrates that have been etched by

the present method infiltrate the voids in the surface region created by removal of the binder phase. The infiltration of the coating is believed to increase the adhesive strength between the coating and the composite material substrate. It has been found that the enhanced adhesion between coatings and composite material substrates achieved by the present method reduces differences in thermal expansion between the substrates and coatings, improves the coatings' resistance to deformation, increases coating wear resistance, and reduces the occurrence of thermal cracking.

As used herein, "composite material" refers to a material, in any form, that includes at least particles of a phase of a hard constituent material and a phase of a binder material that binds together the hard constituent particles. The composite material may be, for example, cemented carbides and cermets. The binder material of the present composite material may include one or a combination of more than one of cobalt, nickel, copper, and iron. In addition to cobalt, nickel, copper, and/or iron, the binder material may include other elements and compounds as are known in the art. Such other elements include, for example, those within Group VIII of the periodic table (elements having atomic numbers 26-28, 44-46, and 76-78), tungsten, zinc, and rhenium. The particles of the hard constituent may be, for example, particles composed of:

- one or more carbide materials selected from tungsten carbide (WC), titanium carbide (TiC), tantalum carbide (TaC), niobium carbide (NbC), vanadium carbide (VC), chromium carbide (Cr_3C_2), molybdenum carbide (MoC), and iron carbide (FeC);

- one or more carbonitrides and/or nitrides of one or more of the refractory metals, including carbonitrides of one or more of W, Ti, Ta, Nb, V, Cr, Mo, and Fe;
- one or more oxides and/or borides of one or more of aluminum, zirconium, and magnesium; and
- one or more of tungsten, molybdenum-based materials, and tungsten-based materials.

As used herein, the term “refractory metals” refers to metals having an extremely high melting point, for example, W, Mo, Ta, Nb, Cr, V, Re, Ti, Pt, and Zr.

In addition to enhancing the adhesion of wear-resistant coatings to the foregoing composite materials, it is believed that the method of the present invention also may be used to enhance the adhesion of wear-resistant and other types of coatings to other types of materials, including, for example, heavy metals, sialons, Si_3N_4 , and composite ceramics, that have a phase that may be etched by the present method. The identities of such other materials may be readily determined by those having ordinary skill in the substrate coating arts. Moreover, although the following examples are directed to the application of wear-resistant coatings to composite material and other substrates, it will be understood that the present method also may be used to better adhere other types of coatings to such substrates. Such other coatings include coatings that impart desirable properties to the substrate surface, such as, for example, coatings that enhance the substrate’s resistance to corrosion, including oxidation, or that provide a particular surface appearance to the substrate. The identities of other coatings that may be applied using the method of the present invention will be readily

apparent to those having ordinary skill in the substrate coating arts once apprised of the invention.

In one embodiment of the method of the present invention, the method generally includes at least the following steps:

- 5 1. Place a composite material substrate to be coated in a chamber of a chemical vapor deposition furnace.
2. Etch away all or a portion of the binder phase in a surface region of the composite material substrate to a depth of about 3 microns to about 15 microns by contacting the surface region with a mixture comprising an etchant gas and an inert gas such as nitrogen gas. (The etchant gas may be selected from, for example, gaseous hydrogen chloride, gaseous H_2F_2 , or the gaseous form of any of the Group VIIA elements. Other suitable etchant gases will be apparent to those of ordinary skill in the art or may be determined by such persons without undue experimentation, and it will be understood that the identity of such suitable alternative etchant gases will depend on the particular composition of the material that is to be etched. The gaseous mixture is applied to the surface of the material to be etched under conditions and for a time suitable to remove the desired amount of binder phase from the material. Such conditions and times may be readily ascertained, without significant experimentation, by those having ordinary skill in the substrate coating arts.)

3. Purge the chamber with a flow of an inert gas ("inert" meaning that it will not react with the binder material) such as, for example, nitrogen, argon, or helium gas.
4. Coat the etched region of the composite material substrate with at least one layer of a wear-resistant material by introducing a reactive gaseous form of the wear-resistant material into the chamber under conditions that will result in the deposition of the wear-resistant material on the etched region. (Such conditions, which generally include such parameters as reactive gas flow rates, chamber gas pressure, chamber and/or substrate temperature, and reaction time) may be readily ascertained by those having ordinary skill in the substrate coating arts once apprised of the present invention.

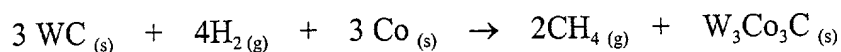
Although the method of the invention is disclosed above as being carried out in a chamber of a chemical vapor deposition (CVD) furnace, it will be understood that the etching step may be carried out in any chamber that is sealed from the environment and into which a flow of the gases may be introduced. An advantage of carrying out the process in a CVD furnace is that the etching, purging, and coating steps may be carried out sequentially in the furnace chamber without the need to move the composite materials from one chamber to another during the process. Thus, the method of the invention may be programmed as a complete cycle in the CVD furnace and accomplished in one run. This feature of the invention provides a distinct advantage over procedures wherein a liquid solution etchant is used to remove binder phase material because such liquid solutions cannot be introduced into the same

chamber employed to coat the substrate by a CVD or PVD process. Also, it has been found that the substrate may be kept cleaner and the depth of etching may be better controlled when using a gaseous etchant as opposed to a liquid etchant.

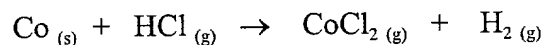
The step of etching binder phase from the composite material substrate preferably should remove binder material to a depth of about 3 to about 15 microns and more preferably about 4 to about 6 microns, into the substrate surface. Too shallow an etching depth does not provide a significant enhancement in coating adhesion. Too great an etching depth weakens the surface of the substrate. Etching time may be varied to account for differences in the susceptibility of the particular binder phase to be removed by the etchant gases. Those having ordinary skill in the substrate coating arts may readily determine the etching time necessary to provide a desired depth of etching for a particular substrate. The substrate temperature at which the etching step should be carried out to remove the desired amount of binder material also will depend upon the character of the binder, but may be readily determined.

Deposits of a phase of W_xCo_yC (wherein $x = 3-9$ and $y = 2-6$), also known as η (eta) phase, may form on the surface of composite material substrates. Eta phase is a hard and brittle carbon-deficient phase that may easily fracture and may be produced when etching substrates that include tungsten, carbon, and cobalt. The presence of eta phase significantly degrades the properties of composite material substrates used in material removal (*i.e.*, cutting, drilling, threading, boring, etc.) applications and, therefore, the generation of eta phase preferably should be avoided during the etching and coating of composite material substrates and other substrates by appropriately adjusting the etching and coating conditions. For example, relative to composite

material substrates including nickel binder, composite material substrates having cobalt binder should be etched at lower substrate temperatures in order to inhibit the formation of eta phase on the surface of the substrate. The inventors also have determined that if hydrogen gas is present during an etching step employing a gaseous etchant, the hydrogen may combine with any carbon present as WC and any cobalt within the substrate material and will thereby make the WC deficient in carbon, resulting in formation of an eta phase. One possible reaction representative of formation of an eta phase is as follows:



The eta phase does not convert to CoCl_2 , as is required for the precursor elements of the eta phase to leave the substrate surface as a gas. The inventors have concluded that formation of eta phase is significantly inhibited when using nitrogen or certain other gases in substitution for hydrogen gas used in conjunction with etchant gas during the etching step. A representation of a possible reaction occurring during etching of a cobalt-containing composite material by a hydrogen chloride etchant gas, and wherein the etchant gas is not applied to the material in combination with hydrogen gas, is believed to be as follows:



The CoCl_2 is a gaseous product that is swept from the coating furnace during the purging step.

Accordingly, the inventors have discovered that the step of etching a substrate including tungsten, carbon, and a binder phase including cobalt will not be satisfactorily accomplished if the gaseous etchant mixture includes hydrogen gas.

For example, when etching binder phase from a composite material cutting insert composed of tungsten carbide particles in a binder composed predominantly of cobalt using a gaseous etchant mixture of hydrogen chloride and hydrogen gases, cobalt residue remains in the voids etched between the tungsten carbide particles and the undesirable eta phase may form, significantly reducing substrate toughness. The inventors have found that nitrogen gas may be advantageously substituted for hydrogen gas to prevent formation of eta phase. More broadly, to better ensure removal of etched binder material so as to avoid formation of eta phase on substrate surfaces, gases that may be substituted for hydrogen gas in the gaseous mixture used in the substrate etching step include those selected from one or more of nitrogen gas and other gases that do not react with the substrate or removed binder and that do not change the oxidation state of the substrate. Such other gases are believed to include, for example, helium, argon, and neon gases.

The foregoing representations of reactions that may occur during the etching process are provided only to better illustrate possible reaction mechanisms, and should not be considered to in any way limit the scope of the invention.

As discussed above, the etchant gas that may be used in the etching step of the method of the present invention may be any gas that will suitably remove the desired depth of binder phase in a surface region of the composite material substrate that is being etched. Such etchant gases include, for example, HCl gas, H_2F_2 gas, and the gaseous form of any of the Group VIIA elements in the periodic table of the elements.

The purging step occurring subsequent to the etching step is necessary to remove any products of the etching reaction and any etchant remaining in the

chamber, and to reduce any explosion hazard. Any gas or combination of gases that will suitably remove the reactant products and remaining etchant gases and that will not react with the binder or hard particle constituents of the composite material may be used as the purging gas. Suitable purging gases include, for example, one or more
5 of nitrogen, helium, and argon gases.

Once etched, the substrate may then be coated with a wear-resistant or other coating material by any conventional composite substrate coating process. Such processes include, for example, CVD, PVD, plasma arc, and super lattice processes. Still other composite material coating procedures will be readily apparent to those of
10 ordinary skill in the substrate coating arts. All such other suitable coating processes may be used in the present method subsequent to the gas etching procedure. Any coating process used to deposit wear-resistant material on a composite material substrate etched by the procedure of the present method is carried out under conditions by which the wear-resistant material may at least partially infiltrate the
15 voids in the composite material created by removal of the binder material. One of ordinary skill may readily determine such conditions without undue experimentation.

On a basic level, the present invention also is directed to a method for removing binder material from a region of a composite material, and the inventive method need not include the subsequent coating step. A composite substrate having a
20 roughened surface may be produced by such a method. Roughened composite substrates may be used in a variety of known applications, including, for example, ball point pen balls, wherein a roughened surface provides enhanced traction. Additionally, substrates may be etched by the present method and then coated at some

later time and/or at a different facility, rather than in a single procedure in which the etching and coating steps are combined. One example of a coating that may be applied in a procedure removed in time from the etching procedure and/or at another facility is a diamond coating.

- 5 Following are actual examples illustrating embodiments of the method of the present invention. The following examples are illustrative examples only, and should not be considered to in any way limit the scope of the present invention.

Example 1

- 10 A Bernex 250 CVD coating furnace was prepared by introducing into the coating chamber of the furnace a 10 l/min (liters/minute) flow of hydrogen gas to establish a 200 mbar hydrogen gas pressure within the chamber. The chamber was then heated to 850°C. A cemented carbide substrate composed of H-91 grade material available from Stellram, LaVergne, Tennessee, was placed in the prepared furnace chamber and the chamber atmosphere was heated to 850°. H-91 grade
- 15 material is composed of 88.5 weight percent tungsten carbide, 11.0 weight percent cobalt, and 0.5 weight percent of a mixture of TiC, TaC, and NbC. The material exhibits a hardness of 89.7 HRA, 14.40 g/cc density, and a transverse rupture strength of approximately 389,000 psi.

- 20 The flow of hydrogen gas was then stopped, and a concurrent flow of 20 l/min nitrogen gas and 1 l/min hydrogen chloride gas was introduced into the chamber to provide a chamber pressure of 800 mbar. Binder was etched to a depth of approximately 5 microns into the substrate's surface by the running the concurrent N₂/HCl gas flow into the chamber for 25 minutes, and then discontinuing the flow of

HCl gas. While maintaining the chamber atmosphere at 850°C, the chamber was then purged for 15 minutes by continuing the 20 l/min flow of N₂ gas while establishing a 60 mbar chamber pressure.

After purging the chamber, and without removing the etched substrate from the chamber, the etched substrate was coated with a moderate temperature milling and turning coating (referred to herein as “MT-milling coating”), which is a multi-layer insert coating consisting of two TiN layers of approximately 1 micron with a TiCN layer of approximately 3 microns disposed between the two TiN layers. The MT-milling coating was deposited on the substrate by introducing into the furnace chamber flows of gases that will produce coatings of TiN, TiCN, and then TiN, in that order, as follows.

Before the coating procedure began, the chamber atmosphere was heated to 920°C and the chamber pressure was reset to 160 mbar. After that pressure was established, a first TiN layer was provided on the substrate by initiating a 9 l/min nitrogen gas flow, increasing the hydrogen gas flow to 14 l/min, and initiating a 2.1 ml/min flow of TiCl₄ gas. The concurrent flows of the three gases were allowed to proceed for 60 minutes while the chamber pressure was maintained at approximately 160 millibars. During the 60-minute period, the furnace temperature was lowered 5-10°C every fifteen minutes so as to be at approximately 895°C at the end of the period.

The interposed TiCN coating was produced by lowering the nitrogen gas flow to 8 l/min, and then resetting chamber pressure to 60 mbar. The TiCl₄ gas flow was then raised to 2.4 ml/min. Once all flows were constant, a flow of CH₃CN gas

generated by vaporizing a 0.3-0.4 ml/min flow of liquid CH₃CN flow was initiated. The concurrent gas flows were continued for 2 hours, during the first hour of which the furnace temperature was reduced to 870°C. At the end of the 2-hour period, the flows of CH₃CN and TiCl₄ gases were discontinued.

- 5 To prepare for deposition of the second TiN layer, the flow of nitrogen gas was discontinued, chamber pressure was set to 500 mbar, hydrogen gas flow was reset to 12 l/min, and furnace temperature was set at 940°C. When that temperature was reached, the pressure was set to 60 mbar, hydrogen gas flow was reset to 10.5 l/min, nitrogen gas flow was reset to 4.5 l/min, and TiCl₄ gas flow was reset to 1.4 ml/min.
- 10 On reaching the target 1.4 l/min TiCl₄ gas flow rate, the flows were continued at temperature for 30 minutes, at which time the pressure was reset to 800 mbar and the gas flows were continued for an additional 30 minutes. The furnace was then purged by shutting off the TiCl₄ gas flow, resetting chamber pressure to 600 mbar, raising hydrogen gas flow to 12 l/min, and lowering nitrogen gas flow to 3.5 l/min. The reset
- 15 gas flows were continued for fifteen minutes. The furnace was then subjected to a cool down procedure.

It was observed that the MT-milling coating infiltrated at least a portion of the voids etched in the substrate's surface.

20

Example 2

The Bernex 250 CVD furnace used in Example 1 was prepared using the procedure described in that example. A cermet substrate composed of SD-5 material, but having the same size and shape as the substrate in Example 1, was placed into the

coating furnace and the furnace atmosphere was heated to 920°C. SD-5 material is a cermet grade material available from Stellram, LaVergne, Tennessee, and is composed of TiCN and Mo₂C particles in a Co/Ni binder. SD-5 material has the approximate elemental composition 45.2 Ti, 22.6 Mo, 10.9 C, 2.3 N, 19.0 Ni, and exhibits the following approximate mechanical properties: 91.8 HRA hardness, 6.30 g/cc density, and a transverse rupture strength of 300,000 psi. After heating the furnace atmosphere to 920°C, the Co/Ni binder was then etched to a depth of 5 microns from the substrate's surface using concurrent flows of hydrogen chloride and nitrogen gases at the flow rates, pressure, and reaction time used in Example 1 above.

The furnace chamber was then purged using a 20 l/min flow of N₂ gas for 15 minutes at a chamber pressure of 60 millibars. The etched composite was then coated with an MT-milling coating using the procedure of Example 1. The MT-milling coating infiltrated the etched voids to a depth of 5 microns ± approximately 1 micron, and with approximately 1 micron of the coating disposed above the substrate's surface.

Example 3

Three Stellram cutting inserts of type SEKN-42-AF4B composed of SD-5 material (as described in Example 2) were first etched and then coated with the MT-milling coating by the following procedure.

A CVD furnace chamber was prepared using the procedure described in Example 1. The SD-5 cutting inserts were then inserted into the furnace chamber and were etched using the procedure of Example 2. Once etched, the inserts were coated with MT-milling coating by the procedure of Example 1. The MT-milling coating

produced on the etched inserts by the foregoing procedure was approximately 5
microns in thickness and the surface TiN layer infiltrated the voids etched in the
inserts' surfaces. Approximately 5 microns of the coating extended above the inserts'
surfaces. Figure 1 is a photomicrograph (2040X) of a prepared cross-section through
the surface of one of the etched and coated SD-5 inserts. The photomicrograph shows
the infiltration of the MT-milling coating into the voids etched in the insert surface.
The infiltration of the coating into the voids increased the adherence of the coating to
the insert and improved the thermal shock resistance of the coating.

The three etched and coated SD-5 inserts and three inserts of the same type
that were unetched and uncoated were inserted at one time into a six-insert Teledyne
(Laverne, Tennessee) HSM-3E4-45 EZ shear cutter that was then installed on a 2
H.P. Bridgeport milling machine and tested under the following milling conditions:

8620 steel at 20-25 Rockwell C hardness
800 surface feet per minute
0.050 inch depth of cut
0.004 - 0.005 inches per tooth (feed rate)
16.5 inch length of cut
2.5 inch width of cut

The SD-5 inserts were pulled and inspected after every two 16.5 inch milling
passes. After the initial two passes, one uncoated SD-5 insert had one thermal crack
started and the five remaining SD-5 inserts did not exhibit thermal cracks. After ten
passes, all three unetched/uncoated SD-5 inserts exhibited one or more thermal cracks
while only one etched and coated SD-5 insert exhibited a single thermal crack.

Milling testing was concluded after 18 passes, at which point each unetched/uncoated
insert exhibited 2-4 thermal cracks on their used edges, while only one thermal crack
existed in one etched and coated insert. The conditions of an edge surface of each of

the three etched/coated SD-5 inserts after 10 and 18 passes are shown in Figures 2a-c and 3a-c, respectively. The used edge conditions of the three unetched/uncoated SD-5 inserts after 10 and 18 passes are shown in Figures 4a-c and 5a-c, respectively.

Example 4

5 Three Stellram SEKN-42-AF4B cutting inserts composed of T-14 material were etched and then coated with an MT-milling coating by the method used in Example 3. T-14 material is a milling grade material available from Stellram (LaVergne, Tennessee) having a nominal composition composed of 70 weight percent tungsten carbide and 20 weight percent of a combination of tantalum carbide,
10 niobium, and titanium carbide. Particles of the foregoing material are bound together by a cobalt binder that is 10 weight percent of the total weight of the material. T-14 material typically exhibits a hardness of 91.20 HRA, 12.43 g/cc density, and an average transverse rupture strength of 296,000 psi.

 The three etched and coated T-14 inserts and three unetched and uncoated T-
15 14 inserts of the same type were inserted at one time into a six-insert HSM-3E4-45 EZ shear cutter, installed on a 2 H.P. Bridgeport milling machine, and tested under the following milling conditions:

 4140 steel at 40-45 Rockwell C hardness
 500 surface feet per minute
20 0.050 inch depth of cut
 0.004 - 0.005 inch per tooth (feed rate)
 14 inch length of cut
 2.5 inch width of cut

25 After four passes under the above conditions, the depth-of-cut regions of the etched and coated T-14 inserts showed no evidence of thermal cracking or deformation as examined under a 40X microscope. After four passes all unetched and

uncoated T-14 inserts exhibited numerous thermal cracks with one insert at the thermal crack breakout stage. As used herein, "thermal crack breakout" is the point at which two or more thermal cracks connect and just before the insert surface fractures. Photomicrographs showing the condition of the depth-of-cut region of the three etched and coated T-14 inserts and the three unetched and uncoated T-14 inserts are shown in Figures 6a-c and 7a-c, respectively. The test demonstrated that using the present invention's method to etch composite substrates with cobalt binder to a 10 micron depth and then coating the substrates provides for a coated insert showing significant edge strength and enhanced resistance to coating/edge delamination, coating spalling, and edge fracture. Results similar to the SD-5 insert milling tests of Example 3 were achieved in that the T-14 substrates etched and coated by the present invention's method provide an increased resistance to thermal cracking of the cobalt-based material.

Example 5

Stellram SEKN-42-AF4B type cutting inserts composed of H-91 grade material were obtained. Half of the H-91 inserts were coated with MT-mill coating in a Bernex 325 furnace using an automated procedure substantially similar to the above-described MT-milling coating procedure so as to provide a layered coating on the inserts composed of approximately 1 micron TiN, approximately 3 microns TiCN, and then 1 micron TiN, all such thicknesses being approximate. The remaining H-91 grade inserts were etched in a Bernex 250 CVD furnace using the procedure of Example 1 and the etched inserts were then MT-milling coated in the furnace by the

procedure described in Example 3. Actual total MT-milling coating thicknesses were determined to be 6.2 microns for the unetched/coated inserts and 5.4 microns for the etched/coated inserts. A photomicrograph of the coated surface of one of the etched and coated H-91 inserts is shown in Figure 8. The figure demonstrates the infiltration of the coating into the etched inserts' surfaces.

Single etched/coated or unetched/coated H-91 inserts were installed on a seven-insert Teledyne HSM-5E4-45 5-inch diameter EZ shear cutter, installed on a 25 H.P. Kearney-Trecker milling machine, and tested under the following milling conditions:

- | | |
|----|---|
| 10 | ASTM A536 (F-33100 unified UNS) nodular cast iron |
| | 875 surface feet per minute |
| | 0.125 inch depth of cut |
| | 0.008 feet per tooth |
| | 20 inch length of cut |
| 15 | 4 inch width of cut |

Each insert was pulled after one pass and examined. After one pass, each of the tested unetched/coated inserts exhibited 6-7 thermal cracks, while the tested inserts that had been etched and coated by the method of the present invention exhibited only a single thermal crack. 30X photomicrographs of an etched/coated H-91 insert after one milling pass and an unetched/coated insert after one milling pass are provided as Figures 9 and 10, respectively.

This example 5 compared inserts composed of identical base materials and identical coatings, with the only significant difference being that the test samples of one set were first etched by the present method and the MT-milling coating had infiltrated the resulting interstices in the inserts' surfaces. The etched and infiltrated

inserts exhibited significantly increased resistance to thermal cracking relative to the unetched coated inserts.

Example 6

5 A heavy metal part containing 90 weight percent tungsten metal particles
suspended in 10 weight percent of an iron/nickel binder was etched and coated with
an MT-milling coating using the procedures of the invention as generally described in
the foregoing examples involving insert composed of SD-5 material. 21X
photomicrographs of sections of the etched and coated metal part taken through the
10 coated surface are shown in Figures 11a-11d. The photomicrographs shown the
infiltration of the coating into the voids etched in the metal part's iron-nickel binder. —

 In each of the foregoing Examples 1-5, all composite material substrates that
were etched using the method of the present invention failed to show evidence of the
15 formation of an eta phase.

 Those of ordinary skill in the substrate coating and treatment arts will
appreciate that various modifications and changes in the details of the invention that
has been disclosed herein may be made without detracting from the advantages
20 provided by the invention, and it will be understood that all such changes and
modifications remain within the principle and scope of the invention as expressed in
the appended claims.

CLAIMS

We claim:

1. A method for removing a portion of the binder phase from a surface of a substrate comprising particles of at least a first phase joined together by the binder phase,
5 the method comprising the act of etching said surface of said substrate by contacting said surface with a gas flow comprising an etchant gas and a second gas for a period of time that will remove said portion of said binder phase, said second gas comprising one or more gases that will not react with said substrate or said portion and that will not change the oxidation state of said substrate during
10 said etching step.
2. The method recited in claim 1 wherein said second gas will not react with said substrate during said etching act to form an eta phase on said surface.
- 15 3. The method recited in claim 2 wherein said second gas is one or more selected from nitrogen gas, helium gas, argon gas, and neon gas.
4. The method recited in claim 3 wherein said etchant gas is one or more selected from hydrogen chloride gas, H_2F_2 gas, and gaseous forms of any of the Group
20 VIIA elements.
5. The method recited in claim 4 wherein said gas flow comprises concurrent flows of hydrogen chloride gas and nitrogen gas.

6. The method of claim 1 wherein in said etching step said binder phase is removed from said surface to a depth of between about 3 microns to about 15 microns.

5 7. The method of claim 6 wherein in said etching step said binder phase is removed from said surface to a depth of between about 4 microns to about 6 microns.

8. The method of claim 1 wherein said substrate is of a composite material, said composite material comprising particles of a hard constituent material and a binder material, said binder material joining together said particles of said hard constituent material.

10

9. The method of claim 8 wherein said composite material is selected from cemented carbides and cermets.

15 10. The method of claim 8 wherein said hard constituent material comprises one or more material selected from the group consisting of:

a carbide material selected from the group consisting of tungsten carbide, titanium carbide, tantalum carbide, niobium carbide, vanadium carbide, chromium carbide, molybdenum carbide, and iron carbide;

20

a carbonitride of a refractory metal;

a nitride of a refractory metal;

a carbonitride of an element selected from the group consisting of W, Ti, Ta, Nb, V, Cr, Mo, and Fe;

an oxide of an element selected from the group consisting of aluminum, zirconium, and magnesium;

5 a boride of an element selected from the group consisting of aluminum, zirconium, and magnesium; and

a material selected from the group consisting of tungsten, a molybdenum-containing material, and a tungsten-containing material.

10 11. The method of claim 8 wherein said binder material comprises one or more materials selected from the group consisting of cobalt, nickel, iron, elements within Group VIII of the periodic table, copper, tungsten, zinc, and rhenium.

12. The method of claim 8 wherein said hard constituent material comprises tungsten
15 carbide and wherein said binder phase comprises cobalt.

13. The method of claim 12 wherein said gas flow comprises hydrogen chloride gas and nitrogen gas.

20 14. The method of claim 1 wherein said etching act occurs within a chamber into which said gas flow is introduced.

15. The method of claim 1 further comprising the act of depositing a coating on said surface of said substrate after said etching act, and wherein at least a portion of said coating infiltrates voids in said surface produced by removal of said binder material from said surface of said substrate.
16. The method of claim 15 wherein said coating enhances the wear resistance of said substrate.
17. The method of claim 16 wherein said coating is comprised of one or more materials selected from the group consisting of TiC, TiN, TiCN, diamond, Al₂O₃, TiAlN, HfN, HfCN, HfC, ZrN, ZrC, ZrCN, BC, Ti₂B, MoS, Cr₃C₂, CrN, CrCN, and CN.
18. The method of claim 16 wherein said coating is an MT-milling coating.
19. The method of claim 1 wherein said substrate is selected from the group consisting of metal cutting inserts, dies, punches, stamps, threading devices, blanking devices, milling devices, turning devices, drilling devices, boring devices, mining bits, drilling bits, tricone bits, percussive bits, road planing devices, wood working bits, wood working blades, drawing devices, heading

devices, back extrusion devices, rod mill roll devices, and wear parts used in corrosive environments.

20. A method for applying a coating at least a portion of the surface of a substrate,
5 the substrate comprising particles of a hard constituent material in a binder phase,
the method comprising the acts of :

removing a portion of said binder phase from said surface of said
substrate by contacting said surface with a gas flow comprising an etchant gas
and a second gas for a period of time that will remove said portion of said binder
10 phase to thereby provide an etched surface on said substrate, said etched surface
comprising voids produced by removal of said portion of said binder phase, said
second gas comprising one or more gases that will not react with said substrate or
said portion and that will not change the oxidation state of said substrate during
said etching step; and

15 applying said coating to at least a portion of said etched surface, at
least a portion of said coating being deposited within at least a portion of said
voids in said etched surface.

21. The method recited in claim 20 wherein said second gas will not react with said
20 substrate or said binder phase removed from said surface during said removing
act to form an eta phase within said voids on said etched surface.

22. The method recited in claim 20 wherein said second gas is one or more selected from nitrogen gas, helium gas, argon gas, and neon gas.

23. The method recited in claim 20 wherein said etchant gas is one or more selected from hydrogen chloride gas, H_2F_2 gas, and gaseous forms of any of the Group VIIA elements.

24. The method recited in claim 23 wherein said gas flow comprises concurrent flows of hydrogen chloride gas and nitrogen gas.

25. The method of claim 20 wherein in said etching step said binder phase is removed from said surface to a depth of between about 3 microns to about 15 microns.

26. The method of claim 25 wherein in said etching step said binder phase is removed from said surface to a depth of between about 4 microns to about 6 microns.

27. The method of claim 20 wherein said substrate is of a composite material, said composite material comprising particles of a hard constituent material and a binder material, said binder material joining together said particles of said hard constituent material.

28. The method of claim 27 wherein said composite material is selected from cemented carbides and cermets.

29. The method of claim 27 wherein said hard constituent comprises one or more material selected from the group consisting of:

a carbide material selected from the group consisting of tungsten carbide,
5 titanium carbide, tantalum carbide, niobium carbide, vanadium carbide,
chromium carbide, molybdenum carbide, and iron carbide;

a carbonitride of a refractory metal;

a nitride of a refractory metal;

a carbonitride of an element selected from the group consisting of W, Ti,
10 Ta, Nb, V, Cr, Mo, and Fe;

an oxide of an element selected from the group consisting of aluminum,
zirconium, and magnesium;

a boride of an element selected from the group consisting of aluminum,
zirconium, and magnesium; and

15 a material selected from the group consisting of tungsten, a
molybdenum-containing material, and a tungsten-containing material.

30. A coated substrate produced by the method of claim 20 wherein said hard
constituent material of said substrate comprises WC, said binder phase comprises
20 cobalt, and said gas flow comprises concurrent flows of hydrogen chloride gas and
nitrogen gas.

31. The method of claim 27 wherein said binder material comprises one or more materials selected from the group consisting of cobalt, nickel, iron, elements within Group VIII of the periodic table, copper, tungsten, zinc, and rhenium.

5 32. The method of claim 27 wherein said hard constituent material comprises tungsten carbide and wherein said binder phase comprises cobalt.

33. The method of claim 20 wherein said act of removing occurs within a chamber into which said gas flow is introduced.

10

34. The method of claim 20 wherein said coating enhances the wear resistance of said substrate.

15

35. The method of claim 34 wherein said coating is comprised of one or more materials selected from the group consisting of TiC, TiN, TiCN, diamond, and Al_2O_3 , TiAlN, HfN, HfCN, HfC, ZrN, ZrC, ZrCN, BC, Ti_2B , MoS, Cr_3C_2 , CrN, CrCN, and CN.

20

36. The method of claim 34 wherein said coating is an MT-milling coating.

37. The method of claim 20 wherein:

said substrate is selected from the group consisting of metal cutting inserts, dies, punches, stamps, threading devices, blanking devices, milling

devices, turning devices, drilling devices, boring devices, mining bits, drilling bits, tricone bits, percussive bits, road planing devices, wood working bits, wood working blades, drawing devices, heading devices, back extrusion devices, rod mill roll devices, and wear parts used in corrosive environments; and

5 said coating enhances the wear resistance of said substrate.

38. The method of claim 37 wherein said coating is an MT-milling coating.

39. An object comprising:

10 a substrate comprised of a composite material, said composite material comprising particles of a hard constituent material and a binder material, said binder material joining together said particles of said hard constituent material, said substrate further comprising a surface portion having voids thereon provided by removing a portion of said binder material from said surface portion, said

15 portion of said binder material being removed from said surface portion by contacting said surface portion with concurrent flows of at least a first gas and a second gas, said first gas capable of removing said binder material from said substrate, said second gas incapable of reacting with said substrate or changing the oxidation state of said substrate during removal of said portion of said binder

20 material; and

 a coating adhered to at least a portion of said surface portion of said substrate, at least a portion of said coating being deposited within at least a portion of said voids, said voids lacking eta phase deposits therein.

40. The object of claim 39 wherein said second gas is one or more selected from nitrogen gas, helium gas, argon gas, and neon gas, and wherein said first gas is one or more selected from hydrogen chloride gas, H_2F_2 gas, and gaseous forms of any of the Group VIIA elements.
41. The object of claim 39 wherein said voids have a depth of between about 3 microns to about 15 microns.
42. The object of claim 39 wherein said composite material is selected from cemented carbides and cermets.
43. The object of claim 39 wherein said hard constituent material comprises one or more material selected from the group consisting of:
- a carbide material selected from the group consisting of tungsten carbide, titanium carbide, tantalum carbide, niobium carbide, vanadium carbide, chromium carbide, molybdenum carbide, and iron carbide;
 - a carbonitride of a refractory metal;
 - a nitride of a refractory metal;
 - a carbonitride of an element selected from the group consisting of W, Ti, Ta, Nb, V, Cr, Mo, and Fe;
 - an oxide of an element selected from the group consisting of aluminum, zirconium, and magnesium;

a boride of an element selected from the group consisting of aluminum, zirconium, and magnesium; and

a material selected from the group consisting of tungsten, a molybdenum-containing material, and a tungsten-containing material.

5

44. The object of claim 39 wherein said hard constituent material of said substrate comprises WC, said binder phase comprises cobalt, said first gas comprises hydrogen chloride gas, and said second gas comprises nitrogen gas.

10 45. The object of claim 39 wherein said binder material comprises one or more materials selected from the group consisting of cobalt, nickel, iron, elements within Group VIII of the periodic table, copper, tungsten, zinc, and rhenium.

15 46. The object of claim 39 wherein said coating enhances the wear resistance of said substrate and is comprised of one or more materials selected from the group consisting of TiC, TiN, TiCN, diamond, Al₂O₃, TiAlN, HfN, HfCN, HfC, ZrN, ZrC, ZrCN, BC, Ti₂B, MoS, Cr₃C₂, CrN, CrCN, and CN.

47. The object of claim 39 wherein said coating is an MT-milling coating.

20

48. The object of claim 39 wherein:

said object is selected from the group consisting of metal cutting inserts, dies, punches, stamps, threading devices, blanking devices, milling

devices, turning devices, drilling devices, boring devices, mining bits, drilling bits, tricone bits, percussive bits, road planing devices, wood working bits, wood working blades, drawing devices, heading devices, back extrusion devices, rod mill roll devices, and wear parts used in corrosive environments; and

5

said coating enhances the wear resistance of said substrate.

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ABSTRACT OF THE INVENTION

Method for removing a portion of the binder phase from the surface of a substrate that is composed of particles of at least a first phase joined together by the binder phase, and wherein the surface is etched by contacting it with a gas flow of an etchant gas and a second gas. The second gas is one or more gases that will not react with the substrate or the removed binder phase and will not alter the oxidation state of the substrate during etching.

10

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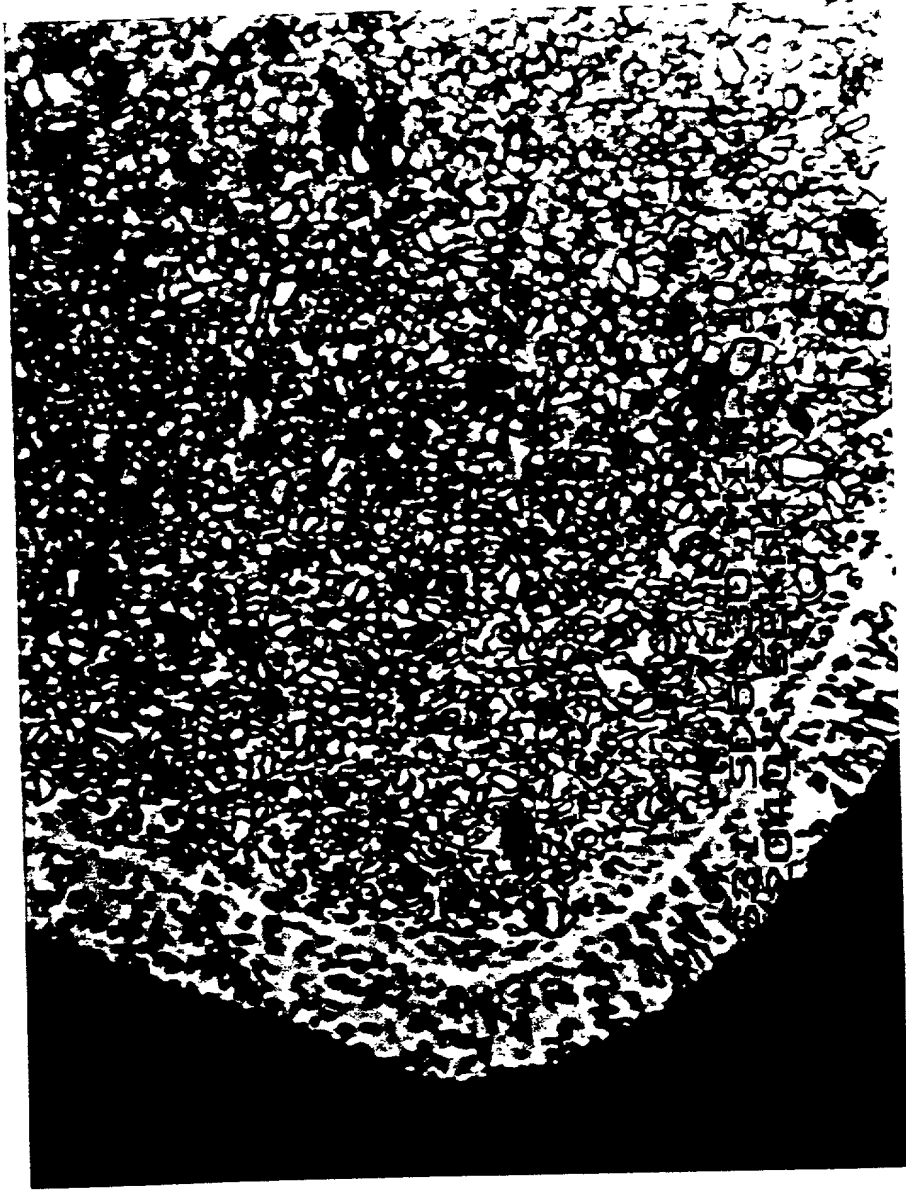


FIG. 1

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FIG. 2a

008240" 10822960



FIG. 2b

008220" F0822960



FIG. 2c

008220" T0822960



FIG. 3a

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FIG. 3b

008220" 70822960



FIG. 3c

008220" 10822960

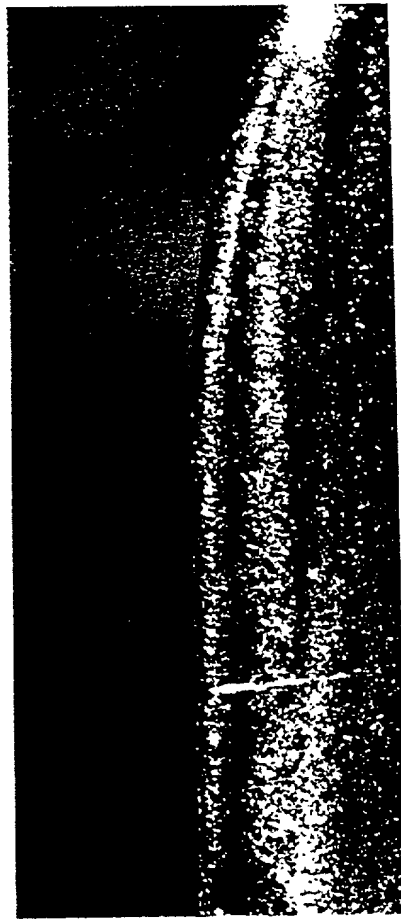


FIG. 4a

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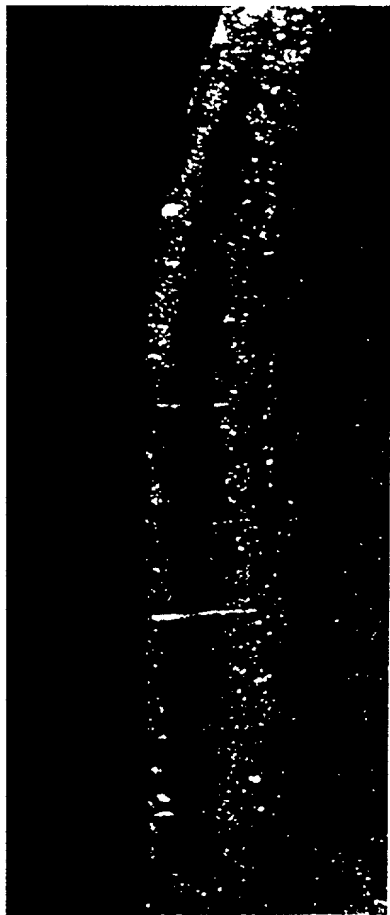


FIG. 4b

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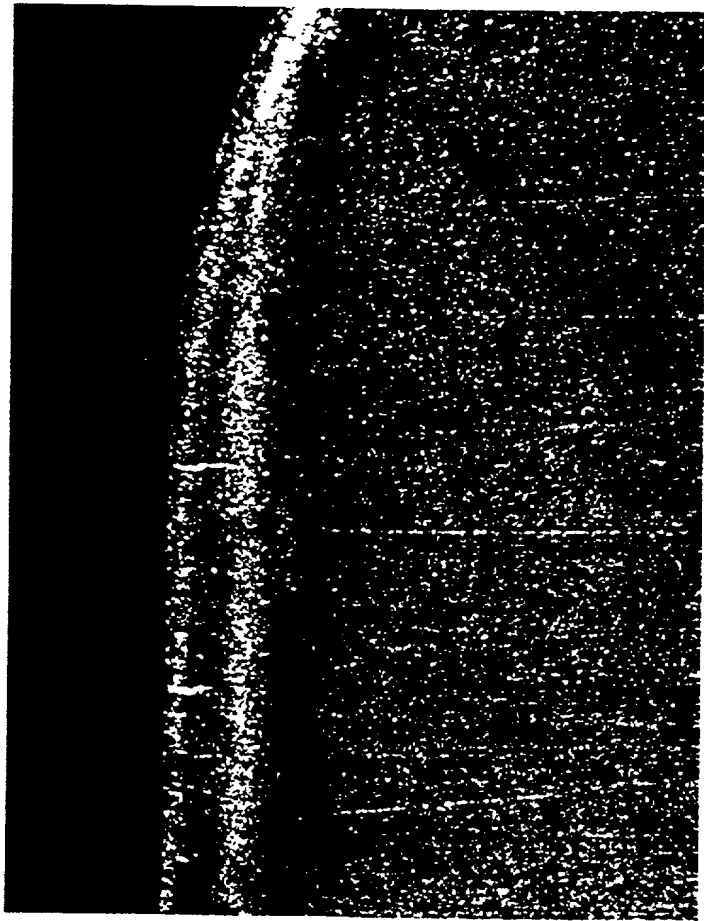


FIG. 4c

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FIG. 5a

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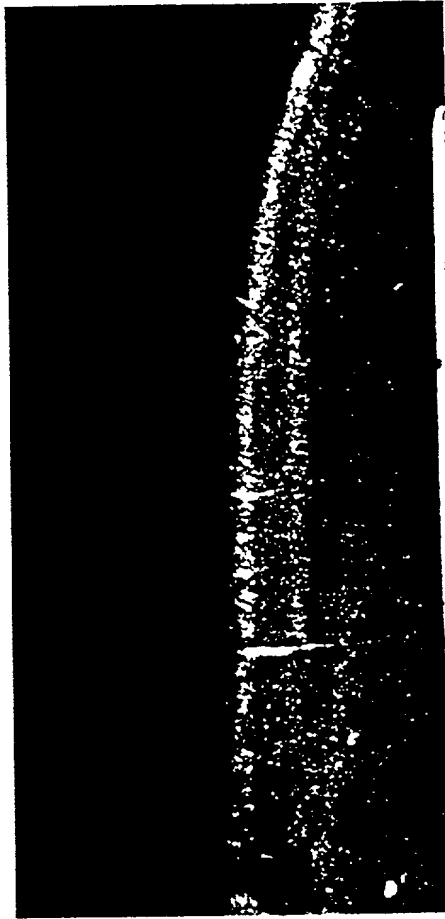


FIG. 5b

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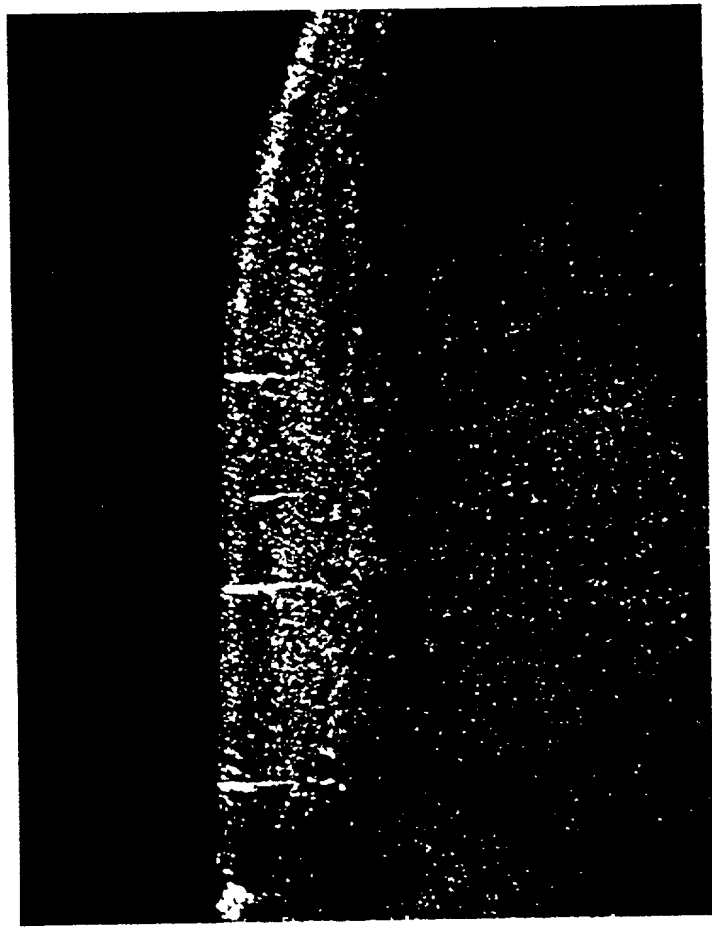


FIG. 5c

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FIG. 6a

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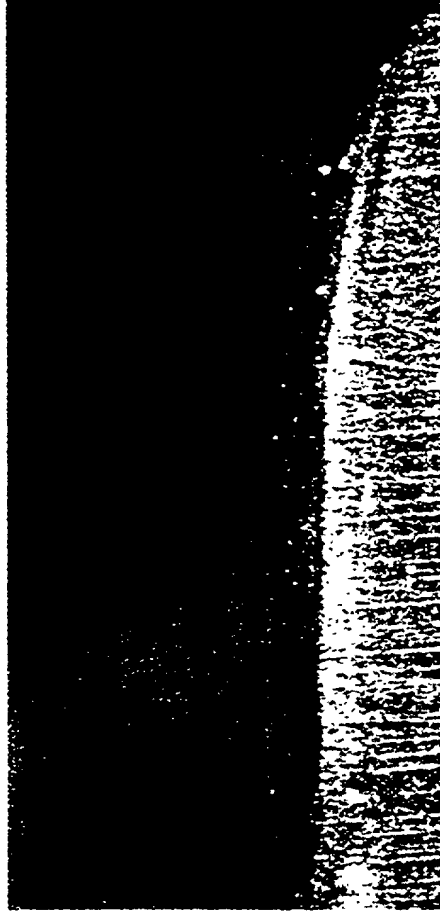


FIG. 6b

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FIG. 6c

09627801.072800



FIG. 7a

09627801.072800



FIG. 7b

008270" T0872960

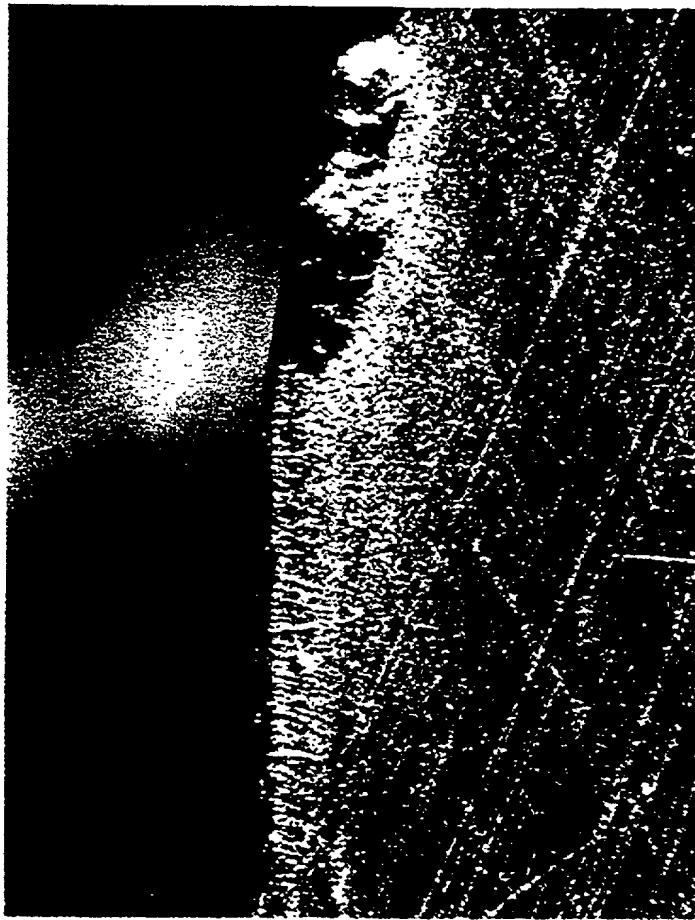


FIG. 7c

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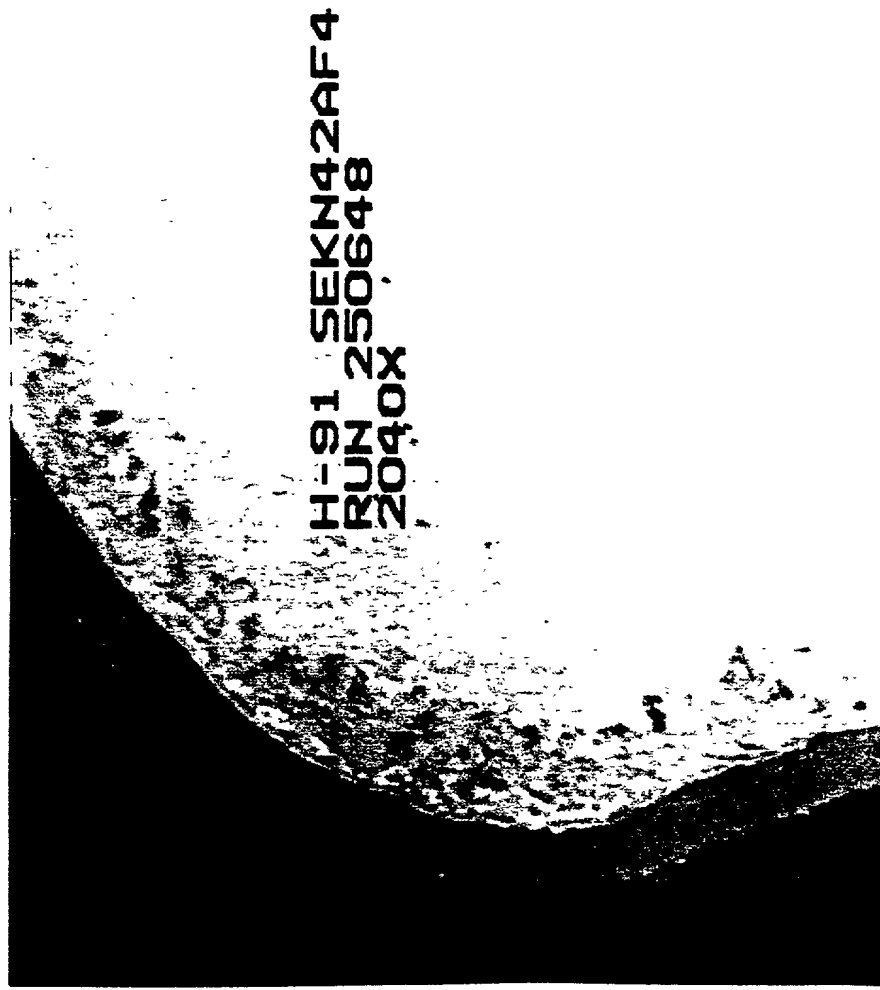


FIG. 8

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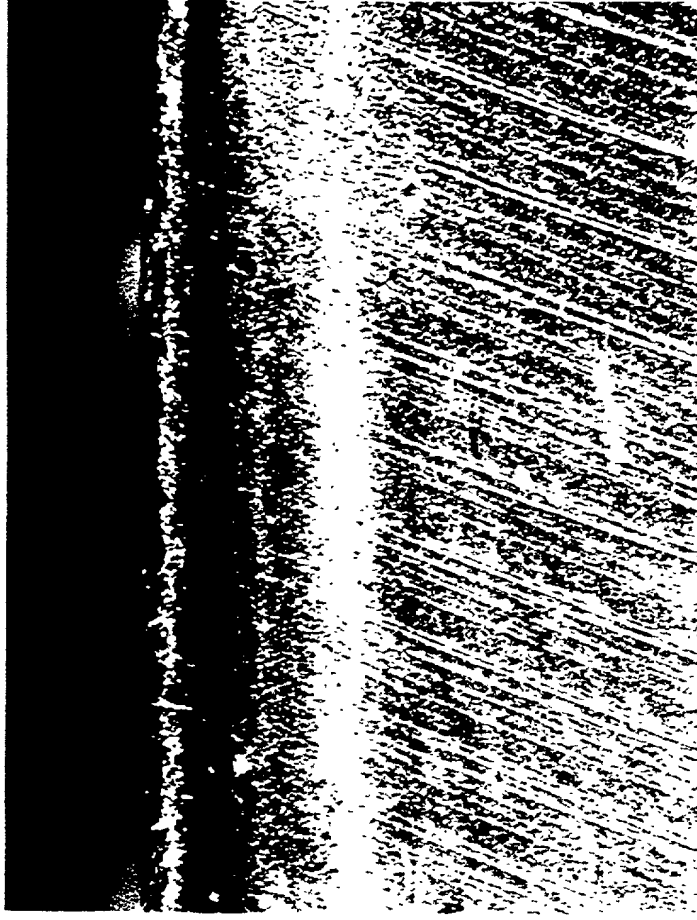


FIG. 9



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FIG.
11a



FIG. 11b

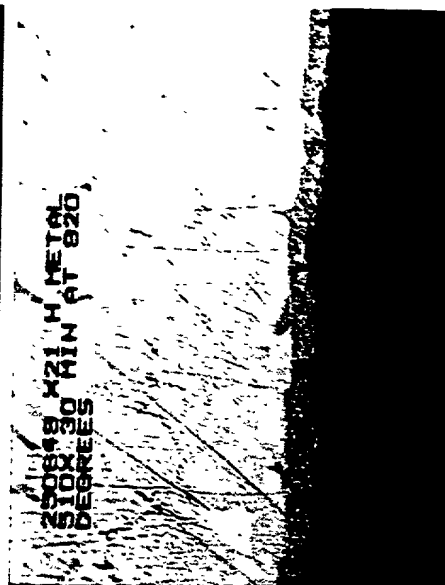


FIG.
11c

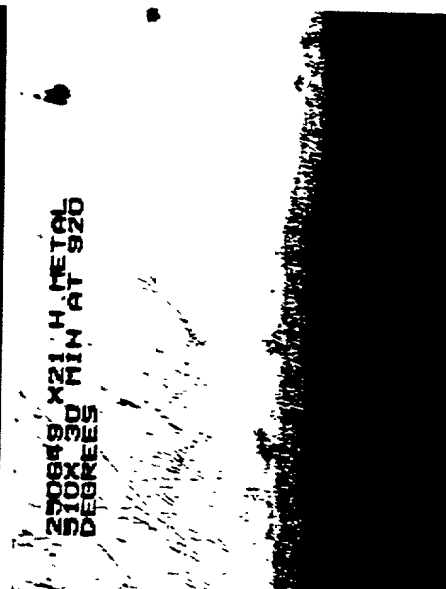


FIG.
11d

DECLARATION

INVENTORSHIP IDENTIFICATION

As a below named inventor, I hereby declare that my residence, post office address and citizenship are as stated below, next to my name. I believe that I am the original, first and sole inventor (*if only one name is listed below*) or an original, first and joint inventor (*if plural names are listed below*) of the subject matter that is claimed, and for which a patent is sought on the invention entitled:

TITLE OF INVENTION

SPECIFICATION IDENTIFICATION

the specification of which:

- (a) ☒ is attached hereto.
- (b) ☐ was filed on _____, as ☐ Serial No. 0/
☐ and was amended on _ (*if applicable*).
- (c) ☐ was described and claimed in PCT International Application No. _ filed on _____
☐ and was amended on _____. (*if applicable*).

ACKNOWLEDGMENT OF REVIEW OF PAPERS AND DUTY OF CANDOR

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims as amended by any amendment referred to above.

I acknowledge the duty to disclose information, which is material to patentability as defined in 37, Code of Federal Regulations, § 1.56.

PRIORITY CLAIMS UNDER 35 U.S.C. § 119(a)-(d)
(Prior Foreign/Pct Application(S) Filed Within 12 Months
(6 Months For Design) Prior To This Application)

I hereby claim foreign priority benefits under Title 35, United States Code, § 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate(s) or 365(a) of any PCT international application(s) which designated at least one country other than the United States of America, listed below and have also identified below any foreign application(s) for patent or inventor's certificate(s) or any PCT international application(s) having a filing date before that of the application(s) of which priority is claimed.

Application Number	Country or PCT	Date Of Filing (Day, Month, Year)	Priority not Claimed	Certified Copy Attached?
			<input type="checkbox"/>	<input type="checkbox"/> YES <input type="checkbox"/> NO

CLAIM FOR BENEFIT OF PRIOR U.S. PROVISIONAL APPLICATION(S) UNDER 35 U.S.C. § 119(e)

I hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below:

Provisional Application Number	Filing Date

CLAIM FOR BENEFIT OF EARLIER US/PCT APPLICATION(S) UNDER 35 U.S.C. 120(All Foreign Application(S), If Any, Filed More Than 12 Months
(6 Months For Design) Prior To This U.S. Application)

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) or 365(c) PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT international application(s) in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37, Code of Federal Regulations, § 1.56, which became available between the filing date of the prior application(s) and the national or PCT international filing date of this application.

U.S Parent Application No.	PCT Parent Application No.	Date Of Filing (Day, Month, Year)	Parent Patent No. (If applicable)

DECLARATION

I hereby declare that my presentation of this paper constitutes a certification under 37 C.F.R § 10.18, which provides, in part, that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and that further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful statements may jeopardize the validity of the application and any patent issuing therefrom.

SIGNATURE(S)**Inventor(s)**

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Inventor's signature

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

POWER OF ATTORNEY

Docket No.
TAM-1629DIV1

Name of Applicant: **TDY Industries, Inc.**
Address of Applicant: **1000 Six PPG Place**
Pittsburgh, PA 15222

Title: **SUBSTRATE TREATMENT METHOD**

Serial No., if Any:

Filed:

TO THE ASSISTANT COMMISSIONER FOR PATENTS

The Assistant Commissioner for Patents
Washington, D.C. 20231

Honorable Sir:
I hereby appoint:

Patrick J. Viccaro, Reg. No. 27,842


Mark Leslie, Reg. No. 36,360

as principal attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith.

Please direct all future correspondence to:

Patrick J. Viccaro
Allegheny Technologies Incorporated
1000 Six PPG Place
Pittsburgh, PA 15222

By:



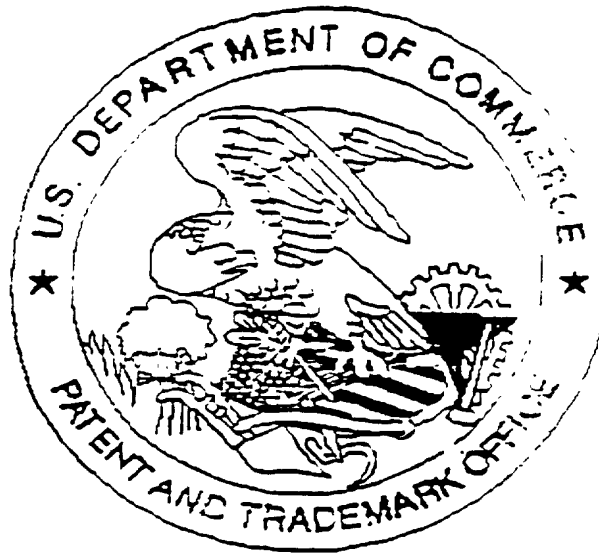
Mary W. Snyder

TDY Industries, Inc.

July 28, 2000

Dated: _____

United States Patent & Trademark Office
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There are 23 sheets of drawings

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